Thermodynamics

1.	Hot Milk in a thermos flask is an example for								
	1) Isolated system	2) Open system	3) Closed system	4) Adiabatic system					
2.	In open system, system and surroundings exchange								
	1) Energy only	2) Matter only 3) Bo	oth 1 & 2 4) Neither	1 nor 2					
3.	Which of the following is a state function?								
	1) Intrinsic energy	2) Enthalpy	3) Heat 4)) Both 1 &2					
4.	The standard heat	The standard heat of combustion of graphite carbon is -393.5 KJ mol ⁻¹ . The							
	standard enthalpy of CO ₂ is								
	1) +393.5 KJ mol ⁻¹	2) -393. 5 KJ mol ⁻¹	3) +196.75 KJ mol ⁻	⁻¹ 4) -196.75 KJ mol ⁻¹					
5.	Which of the follow	ving is a path functio	n?	·					
	1) Internal energy 2) Enthalpy 3) Work 4) Entropy								
6.	Mathematical repr	esentation of 1 st law	v of Thermodynami	ics is					
	1) $Q = E + W$ 2) $H = E + PV$ 3) W	$V = Q \times E$ (4)) ΔH	$\mathbf{H} = \Delta \mathbf{E} + \mathbf{v} \Delta \mathbf{P}$					
7.	Which of the following values of heat of formation indicates that the product is								
	least stable?								
	1) -393.5 KJ	2) -972.7 KJ	3) +89.9 Kj	4) +272.2 KJ					
	Hint: Endothermic	compound is less stab	le. More endothermi	c is least stable.					
8.	Enthalpy change in a cyclic process is								
	1) Infinite	2) Can't be predict	ted 3) Unity	4) Zero					
9.	According to 1st la	w of Thermodynami	cs						
	1) Energy can be created but not destroyed								
	2) Energy cannot be created but can be destroyed								
	3) Energy can be created and destroyed								
	4) Energy neither be created nor destroyed								
10.	Internal energy do	es not include							
	1) Vibrational energ	У	2) Rotational Ener	gy					
	3) Energy due to gravitational pull4) Potential Energy								

11.	At a given temperature internal energy of 4.4gm dry ice is							
	1) Same as 4.4gm liquid CO ₂	2) Same as 4.4gm C	O_2 gas					
	3) Same as 8.8gm dry ice	4) Same as 0.1 moles of dry ice						
12.	The change in internal energy of a sy	stem depends on						
	1) Initial and final states of the system	2) The Path if reve	rsible					
	3) The path if irreversible	4) Initial, final state	es and also on the path					
13.	Enthalpy change during a reaction does not depend upon							
	1) Conditions of a reaction	2) Initial and	d final concentration					
	3) Physical states of reactants and produced	ucts 4) Number o	f steps in the reaction					
14.	The standard enthalpies of n-pentan	ne, isopentane and ne	opentane are -35.0, -					
	37.0 and -40.0 K.cal/mole respective	ly. The most stable	isomer of pentane in					
	terms of energy is							
	1) n-pentane 2) Isopentane	3) Neopentane	4) Both 1 &2					
15.	The enthalpies of the elements in the	ir standard states are	e arbitrarily assumed					
	to be							
	1) Zero at 298 K and 1 atm2) Unity at 298 K and 1 atm							
	3) Zero at all temperatures (4) Zero at 273 K and 1 atm							
16.	The standard enthalpy is zero for the	substance						
	1) C (graphite) 2) C (diamond)	3) CO _{2 (gas)}	4) All					
17.	The heat required to raise the temper	rature of a body by 1^0	C is called					
	1) Specific heat 2) Heat capacity	3) Water equivalent	4) Heat energy					
18.	In exothermic reaction							
	1) $H_R = H_P$ 2) $H_R > H_P$	3) $H_R < H_P$	4) $\Delta H = 0$					
19.	The incorrect IUPAC convention							
1) Heat gained by system +ve sign 2) Work done by system - ve sign								
	3) Work done on the system +ve sign 4) Work done on the system -ve sign							
20.	Which of the following is an endother	rmic reaction?						
	1) $C + O_2 \rightarrow CO_2$ 2) $N_2 + O_2 \rightarrow NO$							
	3) $3H_2 + N_2 \rightarrow 2NH_3$ 4) $PCl_3 + Cl_2 \rightarrow PCl_5$							



1) -RT 2) +RT 3) -2RT 4) Zero **Hint:** $\Delta H = \Delta E + \Delta n RT$ The following is not a combustion reaction 27. 1) $\operatorname{CO} + \frac{1}{2} \operatorname{O}_2 \rightarrow \operatorname{CO}_2$ 2) $\operatorname{C} + \operatorname{O}_2 \rightarrow \operatorname{CO}_2$ 3 $\operatorname{C} + \frac{1}{2} \operatorname{O}_2 \rightarrow \operatorname{CO}_4$) $\operatorname{CH}_4 + 2\operatorname{O}_2 \rightarrow \operatorname{CO}_2 + 2\operatorname{H}_2\operatorname{O}_2$ 28. Match the following. List -I List-II A) $\Delta H < 0$ 1) First law of thermodynamics 2) Heat of combustion **B**) $\Delta H = \Delta E + \Delta nRT$ 3) Relationship between Q_p and Q_v C)Bomb-calorimeter D) Hess law is based on 4) Exothermic reaction **Correct match** С Α B D 3 1. 1 2 2. 4 3 2 3.4 3 1 4. 3 1 $2H_{2(g)} + 2Cl_{2(g)} \rightarrow 4HCl(g), \Delta H^0 = -92.3kJ$ 29. i) If the equation is reversed, the value ΔH^0 equal to +92.3Kj ii) The four H-Cl bonds are stronger than the four bonds in $2H_2$ and $2Cl_2$ iii) The ΔH^0 value will be -92.3Kj if the HCl is produced as a liquid 2. i Only correct 3. i and ii are correct 4. iii Only correct 1. All are correct **30.** The correct statement among the following i) Heat of reaction depends on the temperature at which the reaction is carried ii) ΔH for neutralization is always -Ve.

26. For the reaction $C_2H_{4(g)}+3O_{2(g)}\rightarrow 2CO_{2(g)}+2$ H₂O (g), the difference between

enthalpy change and internal energy change is

iii) Experimentally heat of combustion is ΔE .

1. i only correct 2. ii only correct 3. iii only correct 4. All are correct

31. Match the following.

List-I		List - II
A)solid→vapour		1) 32.8
B) $H_2 + Cl_2 \rightarrow 2HCl$		2) -22
$\Delta H = -44$ K.cals		\frown
heat of formation	n HCl (k.c.	al)
C)Heat of combust	ion of	3) 1 calories
Graphite is-393.5	5 k.J its	
Calorific value o	f (in k.J)	
D) 4.184 Joules is	equal to	4) Endothermic
Correct match is		
A B	С	D
1. 1 2	3	4
2. 4 3	2	1
3.4 2	1	3
4. 2 1	4 C	3
32. Match the following	ıg.	
A) $HNO_3 + KOH$	i) 55.2 l	xJ per mol
В) <i>СН₃СООН</i> + <i>КОН</i>	ii) Path	function
C) Internal Energy	iii) 57.3	kJ per mol
D) Work done	iv) State	function
1) A-(iii), B-(i),	C(iv),	D(ii)
2) A-(i), B-(ii),	C(iii),	D(iv)
3) A-(ii), B-(i),	C(ii),	D(iv)

33. Assertion (A): The enthalpy of formation of $H_2O_{(l)}$ is greater than that of $H_2O_{(g)}$

Reason (**R**): Enthalpy change is negative for the condensation reaction $H_2O_{(g)} \rightarrow H_2O_{(l)}$

1. Both assertion and reason are correct, reason is the correct explanation of the assertion.

2. Both assertion and reason are correct reason is the not correct explanation of the assertion.

List - II

- 3. A is correct, R is incorrect.
- 4. Both A and R are incorrect.
- 34. Match the following.

List-I

- A) Heat of Hydration
- B) Heat of Transition
- C) Molar volume of a gas
- D) Volume of a gas

Is an Intensive property
Is an Extensive property
ΔH is always -Ve
ΔH may be +Ve or -Ve

The correct match is

Α	B	C	D
1) 1	2	3	4
2)4	3	1	2
3) 3	4	51	2
4) 2	1	• 4	3

35. A gas contained a cylinder fitted with a friction less piston expands against a constant pressure 1atm from a volume of 2 liter to volume of 12 liter. In doing so, it absorbs 800J thermal energy from surrounding. Then the △Efor the process is

1) - 213.7 J 2) - 112 J 3) - 50J 4) - 25 J **Solution:** $W = -P \times \Delta V = -1X$ (12-2) =-10lit.atm =-10X101.37kj=-1013.7kj $\Delta E = q - W = 800-1013.7 = -213.7$ Kj

36. 5moles of oxygen are heated at constant volume from 10⁰ to 20⁰C. The change in internal energy of a gas is?

 $\begin{bmatrix} C_P = 7.03 calmol^{-1} des^{-1} and R = 8.31 Jmol^{-1} des^{-1} \end{bmatrix}$

1) 125cal 2) 252cal 3) 50cal 4) 500cal

Solution:

 $C_{P} - C_{V} = R$ $C_{V} = 7.03 - 1.99 = 5.04$

Heat absorbed by 5mole of oxygen in heating from 10 to 20° = 5×*Cr*× ΔT = 5×5.04×10 = 252*cal*.

Since the gas is heated at constant volume, no external work is done W = 0

So change in internal energy will be equal to heat absorbed.

 $\Delta E = q + W = 252 + 0 = 252cal$

37. The amount of work done by 2mole of an ideal gas at 298K in reversible isothermal expansion from 10litre to 20litre is

1) -120J	2) - 2452J	3) -3434.9J	4) 22001
/	/		,

Solution:

 $W = -2.303nRT \log \frac{V_2}{V}$

- = -2.303 X2X8.314 X298Xlog (20/10) =-3434.9j
- 38. 5moles of an ideal gas at 27⁰C expands isothermally and reversibly from a volume of 1L to 10L. The work done in KJ is
 - 1) -14.7 2) -28.72 3) + 28.72 4) 56.72 Solution: $W = -2.303nRT \log \frac{V_2}{V_1}$

-2.303 X5X8.314 X10⁻³ X300Xlog (10/1) =-28.7KJ

- **39. 10 litres of an ideal gas confined to a volume of 10L is released into atmosphere at 300K where the pressure is 1bar. The work done by the gas is** $(R = 0.083L bar K^{-1}mol^{-1})$
 - 1) 249L bar2) 259L bar3) 239L bar4) 220L bar

Solution: $V_2 = \frac{nRT}{P} = 249L$, $W = P\Delta V = 1 \times (249 - 10) = 239L$ bar

40. 1mole of a gas is heated at constant pressure to raise its temperature by 1⁰C. The work done in Joules is

1) -4.3 2) -8.314 3) -16.62 4) Unpredictable Solution: $W=-nR\Delta T = -1 \times 8.314 \times 1 = -8.314J$

41. 3.0 moles of ideal gas is heated at constant pressure from 47⁰C to 147⁰C. Then the work expansion of gas is

1) - 2.494KJ 2) + 2.494KJ 3) - 10.5KJ 4) + 10.5KJ Solution: W=-nR Δ T =-3X 8.314X10⁻³(147-47)=-2.494KJ

- 42. The pKa values of four acids A, B, C and D are 9.14, 9.92, 2.86 and 1.3 respectively. The heat of neutralization is more in the following reaction
 - 1) $A + NaOH \rightarrow \dots$ 2) $B + NaOH \rightarrow \dots$
 - 3) $D + NaOH \rightarrow \dots$ 4) $C + NaOH \rightarrow \dots$

Solution: Lower P_a^k represents strong acid. The heat of neutralization is more for a strong acid.

43. The heats of neutralisation of acids A, B, C and D with NaOH are -13.5 K.cal, -12.7 K.cal, -11.8 K.cal, -12.4 K.cal respectively.

The weakest acid is

1) A 2) B 3) C 4) D

Solution: If the heat of neutralisation is lowest then the acid is weakest.

44. According to $H_2(g) + I_2(g) \rightarrow 2HI(g), \Delta H = 51.9$ KJ. heat of formation of HI is 1) 51.9 KJ 2) -51.9 KJ 3) -25. 95 KJ 4) 25.95 KJ

Solution: heat of formation = Δ H per mole= (51.9/2)=25.95

45. The heat of formation of $H_2O_{(1)}$ is -286.2 KJ. The heat of formation of $H_2O_{(g)}$ is likely to be

- 1) -286.2 KJ 2) -290.78 KJ
- 3) -335.2 KJ 4) -242.76 KJ

Solution: $H_2O_{(1)} \rightarrow H_2O_{(g)}$ is an Endothermic process.

46. $\text{NH}_4\text{Cl}(s) + \text{H}_2\text{O} \rightarrow \text{NH}_4\text{Cl}(aq) \quad \Delta H = 16.3 \text{ KJ}$

$\Delta \mathbf{H}$ in the above reaction represents

- 1) Heat of solution 2) Heat of hydration
- 3) Heat of dilution 4) Heat of ionization

47. A system absorbs 10 kJ of heat and does 4 kJ of work. The internal energy of the system would?

- 1) Decreases by 6 kJ2) Increases by 6 kJ
- 3) Decreases by 14 kJ 4) Decreases by 14 kJ

Solution: q=10Kj, W=-4Kj thus $\Delta E=q+W=10$ -4=6Kj

48. When 4 grams of methane is completely burnt in oxygen, the heat evolved is 224 kJ. What is the heat of combustion (in KJ) of methane?

1) -1120 2) -968 3) -896 4) -560

Solution: Heat of Combustion= heat liberated per 1 mole

if 4gm of methane given 224KJ then 1mole i.e 16gm of methane gives (16/4)x224=896KJ.

49. One mole of ideal gas expands freely at 310 K from five liter volume to 10 liter volume. Then ∆E and∆ H of the process are respectively

1) 0 and 5 cal 2) 0 and 5 x 300 cal 3) 0 and 0 4) 5 and 0 cal **Hint:** for an Ideal gas Δ H=o and Δ E=0

50. The heat of dissociation (in K.cals/mole) of CH₄ and C₂H₆ are 360 and 620 respectively. From these the C - C bond energy in the ethane can be evaluated as

1) 260 2) 130 3) 80 4) 200

Solution:

Average energy of C-H bond in $CH_4=(360/4)=90$

 C_2H_6 has 6 C-H bonds and 1 C-C bond.

6C-Hbonds +1 C-C bond = 540

Energy of C-C = 620-(6C-H) =620-6X90=620-540 = 80K.cal

KEY

1)1	2)3	3)4	4)2	5)3	6)1	7)4	8)4	9)4	10)3
11)4	12)1	13)4	14)3	15)1	16)1	17)1	18)2	19)4	20)2
21)1	22)1	23)1	24)2	25)3	26)4	27)3	28)2	29)3	30)4
31)3	32)1	33)1	34)2	35)1	36)2	37)3	38)2	39)3	40)2
41)1	42)3	43)3	44)4	45)4	46)1	47)2	48)3	49)3	50)3
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